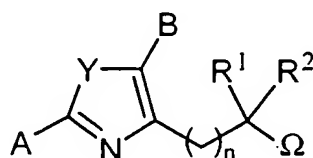


In the Claims:

Claim 1 (currently amended) A method of treating disorders of the central or peripheral nervous system in warm-blooded animals comprising administering to warm-blooded animals in need thereof an amount of a compound of the formula

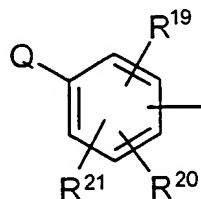


(I)₃

in racemic, enantiomeric form or any combination of these forms,

in which

A is



in which Q is selected from the group consisting of H, -OR²², -NR²³R²⁴, phenyl optionally substituted by at least one substituent independently selected from the group consisting of OH, cyano, nitro, alkyl, alkoxy or -NR¹⁰R¹¹ and a group with two

substituents representing together a methylenedioxy or ethylenedioxy, or Q is selected from the group consisting of -COPh, -SO₂Ph or -CH₂Ph, said COPh, -SO₂Ph or -CH₂Ph optionally substituted on its aromatic parts by at least one independently alkyl or alkoxy or halogen,

R¹⁰ and R¹¹ are independently selected from the group consisting of hydrogen, alkyl and -COR¹², or R¹⁰ and R¹¹ form together with the nitrogen atom an optionally substituted heterocycle containing 4 to 7 members and 1 to 3 heteroatoms including the nitrogen atom already present, the additional heteroatoms being independently selected from the group consisting of O, N and S,

R¹² is selected from the group consisting of hydrogen, alkyl, alkoxy and NR¹³R¹⁴,

R¹³ and R¹⁴ are independently, hydrogen or alkyl, or R¹³ and R¹⁴ form together with the nitrogen atom an optionally substituted heterocycle containing 4 to 7 members and 1 to 3 heteroatoms including the nitrogen atom already present, the additional heteroatoms being chosen independently from the group consisting of O, N and S,

R²² is selected from the group consisting of hydrogen, alkyl and aryl optionally substituted by at least one substituent selected from the group consisting of alkyl, OH, halogen, nitro and alkoxy,

R²³ and R²⁴ are independently selected from the group consisting of hydrogen, alkyl and -CO-R²⁵,

R²⁵ is alkyl,

and R¹⁹, R²⁰ and R²¹ are independently selected from the group consisting of hydrogen, halogen, -OH, -SR²⁶, alkyl, cycloalkyl, alkenyl, alkoxy, cyano, nitro, -SO₂NHR⁴⁹, -CONHR⁵⁵, -S(O)_qR⁵⁶, -NH(CO)R⁵⁷, -CF₃, -OCF₃ and NR²⁷R²⁸,

R^{27} and R^{28} are independently selected from the group consisting of hydrogen, alkyl and $-\text{COR}^{29}$, or R^{27} and R^{28} form together with the nitrogen atom an optionally substituted heterocycle containing 4 to 7 members and 1 to 3 heteroatoms including the nitrogen atom already present, the additional heteroatoms being independently selected from the group consisting of O, N and S,

R^{49} and R^{55} are independently each time that they occur, hydrogen or alkyl or alkylcarbonyl,

q is an integer from 0 to 2,

R^{56} and R^{57} are, each time that they occur, hydrogen or alkyl or alkoxy,

R^{29} is selected from the group consisting of hydrogen, alkyl, alkoxy and $-\text{NR}^{30}\text{R}^{31}$,

R^{30} and R^{31} are independently selected from the group consisting of hydrogen and alkyl, or R^{30} and R^{31} form together with the nitrogen atom an optionally substituted heterocycle containing 4 to 7 members and 1 to 3 heteroatoms including the nitrogen atom already present, the additional heteroatoms being independently selected from the group consisting of O, N and S,

Y is O or S;

R^1 is hydrogen, alkyl, aminoalkyl, alkoxyalkyl, cycloalkyl, cycloalkylalkyl, trifluoromethylalkyl, alkenyl, allenyl, allenylalkyl, alkynyl, cyanoalkyl, $-(\text{CH}_2)_g-\text{Z}^1\text{R}^{39}$, $-(\text{CH}_2)_g-\text{COR}^{40}$, $-(\text{CH}_2)_g-\text{NHCOR}^{70}$, aryl, aralkyl, arylcarbonyl, heteroarylalkyl and aralkylcarbonyl, the aryl group of the aryl, aralkyl, arylcarbonyl, heteroarylalkyl or aralkylcarbonyl itself being optionally substituted by at least one

substituent selected from the group consisting of alkyl, halogen, alkoxy, nitro, cyano, cyanoalkyl, amino, alkylamino, dialkylamino, $-(CH_2)_k-Z^2R^{39}$ and $-(CH_2)_k-COR^{40}$, Z^1 and Z^2 is selected from the group consisting of a bond, -O-, $-NR^{41}$ - and -S-,

R^{39} and R^{41} are independently each time that they occur, selected from the group consisting of hydrogen, alkyl, alkenyl, alkynyl and cyanoalkyl,

R^{40} is, independently each time that it occurs, selected from the group consisting of hydrogen, alkyl, allenyl, allenylalkyl, alkenyl, alkynyl, cyanoalkyl, alkoxy and $NR^{42}R^{43}$,

R^{42} and R^{43} are independently, each time that they occur, selected from the group consisting of hydrogen, alkyl, allenyl, allenylalkyl, alkenyl, alkynyl and cyanoalkyl and R^2 is selected from the group consisting of hydrogen, alkyl, aminoalkyl, alkoxyalkyl, cycloalkyl, cycloalkylalkyl, trifluoromethylalkyl,

$-(CH_2)_g-NHCOR^{71}$, aralkyl and heteroarylalkyl optionally substituted on the aryl or heteroaryl by at least one member selected from the group consisting of halogen, alkyl, alkoxy, hydroxyl, cyano, nitro, amino, alkylamino and dialkylamino,

R^{70} and R^{71} are independently alkyl or alkoxy;

or R^1 and R^2 , taken together with the carbon atom which carries them, form a carbocycle with 3 to 7 members;

B is selected from the group consisting of hydrogen, alkyl, $-(CH_2)_g-Z^3R^{44}$ and carbocyclic aryl optionally substituted 1 to 3 times by a member selected from the group

consisting of halogen, alkyl and alkoxy of 1 to 6 carbon atoms, hydroxy, cyano, nitro, amino, alkylamino, dialkylamino and a carbocyclic aryl,

Z^3 is selected from the group consisting of a bond, -O-, NR^{45} - and -S-,

R^{44} and R^{45} are independently selected from the group consisting of hydrogen, alkyl, alkenyl, alkynyl, allenyl, allenylalkyl and cyanoalkyl;

Ω is $NR^{46}R^{47}$ or OR^{48} ;

R^{46} and R^{47} are independently selected from the group consisting of hydrogen, alkyl, cycloalkyl, cycloalkylalkyl, alkenyl, alkynyl, allenyl, allenylalkyl, cyanoalkyl, $-(CH_2)_g-Z^4R^{50}$, $-(CH_2)_k-COR^{51}$, $-(CH_2)_k-COOR^{51}$, $-(CH_2)_k-CONHR^{51}$, $-SO_2R^{51}$, aryl, aralkyl, aryloxyalkyl, arylcarbonyl, arylimino, aralkylcarbonyl, heteroaryl, the aryl or heteroaryl of said aryl, aralkyl, aryloxyalkyl, arylcarbonyl, arylimino, aralkylcarbonyl, heteroaryl, pyridinylalkyl or pyridinylcarbonyl being optionally substituted by at least one member independently selected from the group consisting of halogen, alkyl, alkoxy, hydroxy, nitro, cyano, cyanoalkyl, amino, alkylamino, dialkylamino, Z^5R^{50} , $-(CH_2)_k-COR^{51}$ and $-(CH_2)_k-COOR^{51}$,

Z^4 and Z^5 are selected from the group consisting of a bond, -O-, $-NR^{52}$ - and -S-, or R^{46} and R^{47} taken together form with the nitrogen atom a non-aromatic heterocycle with 4 to 8 ring members, the elements of the chain being selected from the group consisting of $-CH(R^{53})-$, $-NR^{54}-$, -O-, -S- and -CO-,

R^{50} and R^{52} are independently each time that they occur, selected from the group consisting of hydrogen, alkyl, alkenyl, alkynyl, allenyl, allenylalkyl and cyanoalkyl, R^{51} is, independently each time that they occur, selected from the group consisting of hydrogen, cycloalkyl and cycloalkylalkyl in which the cycloalkyl has 3 to 7 carbon atoms, alkyl of 1 to 8 carbon atoms, alkenyl, alkynyl, allenyl, allenylalkyl, cyanoalkyl, alkoxyalkyl, $NR^{58}R^{59}$, aryl or aralkyl, said aryl or aralkyl optionally substituted by at least one member selected from the group consisting of halogen, alkyl and alkoxy, R^{58} and R^{59} are independently a member selected from the group consisting of hydrogen, alkyl, alkenyl, alkynyl, allenyl, allenylalkyl and cyanoalkyl, R^{53} and R^{54} are independently a member selected from the group consisting of hydrogen or $-(CH_2)_k-Z^7R^{60}$ and $-(CH_2)_k-COR^{61}$, Z^7 is a member selected from the group consisting of a bond, $-O-$, $-NR^{62}-$ and $-S-$, R^{60} and R^{62} are independently a member selected from the group consisting of hydrogen, alkyl, alkenyl, allenyl, allenylalkyl, alkynyl, cyanoalkyl, aryl, aralkyl, arylcarbonyl, aralkylcarbonyl, pyridinyl, pyridinylalkyl or pyridinylcarbonyl, the aryl or pyridinyl of the aryl, aralkyl, arylcarbonyl, aralkylcarbonyl, pyridinyl, pyridinylalkyl or pyridinylcarbonyl being optionally substituted by at least one substituent selected from the group consisting of alkyl, halogen, nitro, alkoxy, cyano, cyanoalkyl, $-(CH_2)_k-Z^8R^{63}$ and $-(CH_2)_k-COR^{64}$, R^{61} is a member selected from the group consisting of hydrogen, alkyl, allenyl, allenylalkyl, alkenyl, alkynyl, cyanoalkyl, alkoxy and $NR^{65}R^{66}$, R^{65} and R^{66} are independently selected from the group consisting of hydrogen, alkyl, allenyl, allenylalkyl, alkenyl, alkynyl and cyanoalkyl,

Z^8 is a member selected from the group consisting of a bond, -O-, -NR⁶⁷- and -S-,

R⁶³ and R⁶⁷ are independently a member selected from the group consisting of hydrogen, alkyl, allenyl, allenylalkyl, alkenyl, alkynyl and cyanoalkyl,

R⁶⁴ is a member of the group consisting of hydrogen, alkyl, allenylalkyl, alkenyl, ~~alkenyl~~, alkynyl, cyanoalkyl, alkoxy and NR⁶⁸R⁶⁹,

R⁶⁸ and R⁶⁹ are independently a member selected from the group consisting of hydrogen, alkyl, allenyl, allenylalkyl, alkenyl, alkynyl cyanoalkyl,

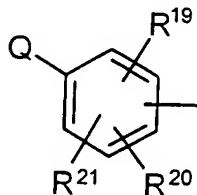
and R⁴⁸ is a member selected from the group consisting of hydrogen, alkyl, alkynyl and cyanoalkyl;

g and p, each time that they occur, being independently integers from 1 to 6, and k and n, each time that they occur, being independently integers from 0 to 6;

or a salt thereof sufficient to inhibit monoamine oxydases and lipidic peroxidation and a modulating activity vis-à-vis sodium channels.

Claim 2 (previously presented) The method of claim 1 wherein

A is



in which Q is selected from the group consisting of H, -OR²², -SR²², -NR²³R²⁴, phenyl optionally substituted by at least one member selected from the group consisting of halogen, OH, cyano, nitro, alkyl, alkoxy and -NR¹⁰R¹¹ and a group of two substituents together being methylenedioxy or ethylenedioxy, or Q is selected from the group consisting of -COPh, -OPh, -SPh, -SO₂Ph or -CH₂Ph, said -COPh, -OPh, -SPh, -SO₂Ph or -CH₂Ph being optionally substituted on its aromatic part by at least one member selected from the group consisting of alkyl, alkoxy and halogen, R¹⁰ and R¹¹ are independently selected from the group consisting of hydrogen, alkyl, or R¹⁰ and R¹¹ form together with the nitrogen atom an optionally substituted heterocycle containing 4 to 7 ring members and 1 to 3 heteroatoms including the nitrogen atom already present, the additional heteroatoms being independently selected from the group consisting of O, N and S, R²² is selected from the group consisting of hydrogen, alkyl and aryl optionally substituted by at least one member selected from the group consisting of alkyl, OH, halogen, nitro and alkoxy, R²³ and R²⁴ are, independently, hydrogen, or alkyl or -CO-R²⁵, R²⁵ is alkyl, and R¹⁹, R²⁰ and R²¹ are independently selected from the group consisting of hydrogen, halogen, OH, SR²⁶, alkyl, cycloalkyl, alkenyl, alkoxy, cyano, nitro, -SO₂NHR⁴⁹, -CONHR⁵⁵, -S(O)_qR⁵⁶, -NH(CO)R⁵⁷, -CF₃, -OCF₃ and NR²⁷R²⁸, R²⁶ is hydrogen or alkyl, R²⁷ and R²⁸ are independently selected from the group consisting of hydrogen, alkyl and

-COR²⁹, or R²⁷ and R²⁸ form together with the nitrogen atom an optionally substituted heterocycle containing 4 to 7 ring members and 1 to 3 heteroatoms including the nitrogen atom already present, the additional heteroatoms being selected from the group consisting of O, N and S,

R⁴⁹ and R⁵⁵ are, independently each time that they occur, hydrogen or alkyl or alkylcarbonyl,

q is an integer from 0 to 2,

R⁵⁶ and R⁵⁷ are, independently each time that they occur, hydrogen or alkyl or alkoxy,

R²⁹ is selected from the group consisting of hydrogen, alkyl, alkoxy and -NR³⁰R³¹,

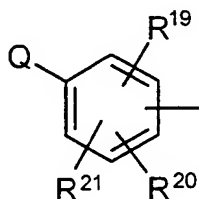
R³⁰ and R³¹ are independently selected from the group consisting of hydrogen, alkyl, or

R³⁰ and R³¹ form together with the nitrogen atom an optionally substituted heterocycle containing 4 to 7 ring members and 1 to 3 heteroatoms including the nitrogen atom already present, the additional heteroatoms being selected from the group consisting of O, N and S.

Claim 3 (previously presented)

The method of claim 1 wherein

A is



in which Q is -OR²², -SR²², -NR²³R²⁴, phenyl optionally substituted by at least one member selected from the group consisting of halogen, OH, cyano, nitro, alkyl, alkoxy and -NR¹⁰R¹¹,

R¹⁰ and R¹¹ are independently selected from the group consisting of hydrogen or alkyl, or R¹⁰ and R¹¹ form together with the nitrogen atom an optionally substituted heterocycle containing 4 to 7 ring members and 1 to 3 heteroatoms including the nitrogen atom already present, the additional heteroatoms being selected from the group consisting of O, N and S,

R²² is selected from the group consisting of hydrogen, alkyl and aryl optionally substituted by at least one member selected from the group consisting of alkyl, OH, halogen, nitro and alkoxy,

R²³ and R²⁴ are, independently, hydrogen or alkyl,

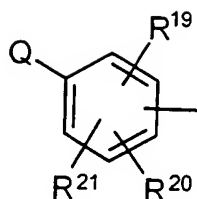
and R¹⁹, R²⁰ and R²¹ are independently selected from the group consisting of hydrogen, halogen, OH, -SR²⁶, alkyl, alkenyl, alkoxy and NR²⁷R²⁸,

R²⁶ is hydrogen or alkyl,

R²⁷ and R²⁸ are independently selected from the group consisting of hydrogen, alkyl, or R²⁷ and R²⁸ form together with the nitrogen atom an optionally substituted heterocycle containing 4 to 7 members and 1 to 3 heteroatoms including the nitrogen atom already present, the additional heteroatoms being selected from the group consisting of O, N and S.

Claim 4 (currently amendment) The method of claim 3 wherein

A is



in which Q is selected from the group consisting of -OR²², -SR²² and phenyl substituted by OH and optionally by at least one member selected from the group consisting of halogen, OH, alkyl and alkoxy,

R²² is hydrogen or alkyl,

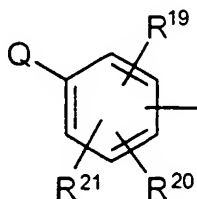
and R¹⁹, R²⁰ and R²¹ are independently selected from the group consisting of hydrogen, halogen, OH, SR²⁶, alkyl and alkoxy,

R²⁶ is hydrogen or alkyl.

Claim 5 (previously presented) The method of claim 4 wherein

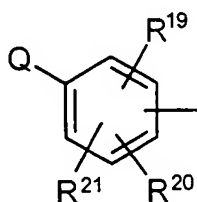
the compound corresponds to formula (I)₃ in which Y is O

A is



in which Q is OH, two of the R^{19} , R^{20} and R^{21} are selected from the group consisting of alkyl, alkoxy, alkylthio, amino, alkylamino and dialkylamino and the third is selected from the group consisting of hydrogen, alkyl, alkoxy, alkylthio, amino, alkylamino and dialkylamino, or Q is phenyl substituted by OH and at least one member selected from the group consisting of halogen, OH, alkyl, alkoxy and $-NR^{10}R^{11}$, R^{10} and R^{11} are independently hydrogen or alkyl.

Claim 6 (previously presented) The method of claim 5 wherein A is

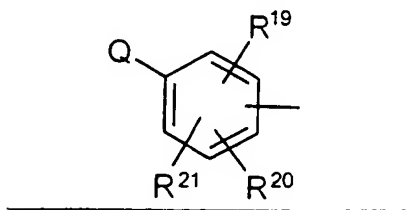


in which Q is OH, two of the R^{19} , R^{20} and R^{21} are alkyl and the third is H,
or Q is phenyl substituted by OH and at least one alkyl.

Claims 7-13 (cancelled)

Claim 14 (currently amended) The method of claim 1 wherein ~~He~~ is
~~formula (I)3 in which~~ Y is O;

A is



in which Q is OH, two of the R^{19} , R^{20} and R^{21} are alkyl and the third is hydrogen,

or in which Q is phenyl substituted by OH and at least one alkyl;

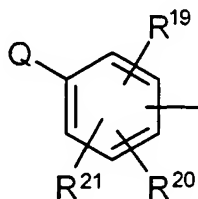
B is hydrogen;

n is 0 or 1;

R^1 and R^2 both are hydrogen;

and Ω is $NR^{46}R^{47}$, R^{46} is selected from the group consisting of hydrogen, alkyl, alkynyl, hydroxyalkyl and cyanoalkyl and R^{47} is hydrogen or alkyl or R^{46} and R^{47} form together with the nitrogen atom which carries them a non-aromatic heterocycle with 5 to 7 ring members, the additional members are $-CH_2-$ or $-NH-$.

Claim 15 (previously presented) The method of claim 14, wherein the compound is selected from the group consisting of



- 4-[3,5-bis(1,1-dimethylethyl)-4-hydroxyphenyl]-N-methyl-2-thiazolemethanamine;
- 2-[(4-[3,5-di(tert-butyl)-4-hydroxyphenyl]-1,3-thiazol-2-yl)methyl](methyl)amino]-acetonitrile;
- 2,6-di(tert-butyl)-4-(2-[(2-hydroxyethyl)(methyl)amino]methyl)-1,3-thiazol-4-yl)phenol;

- 4-[[4-(3,5-ditert-butyl-4-hydroxyphenyl)-1,3-thiazol-2-yl]methyl](methylamino)-butanenitrile;
- 2,6-ditert-butyl-4-(4-{2-[methyl(2-propynyl)amino]ethyl}-1,3-oxazol-2-yl)phenol;
- 3-[[2-[2-(3,5-ditert-butyl-4-hydroxyphenyl)-1,3-oxazol-4-yl]ethyl](methylamino)-propanenitrile;
- 2,6-ditert-butyl-4-{4-[2-(1-piperazinyl)ethyl]-1,3-oxazol-2-yl}phenol;

and the pharmaceutically acceptable salts of the latter.

Claims 16-21 (cancelled).

Claim 22 (currently amended) The method of claim 1 wherein the compound is selected from the group consisting of 2-[3,5-bis(1,1-dimethylethyl)-4-hydroxyphenyl]-4-oxazoleethanol, 2,6-ditert-butyl-4-(4-{2-[methyl(2-propynyl)amino]ethyl}-1,3-oxazol-2-yl)phenol, [{2-[2-(3,5-ditert-butyl-4-hydroxyphenyl)-1,3-oxazol-4-yl]ethyl}(methylamino)]acetonitrile, 3-[[2-[2-(3,5-ditert-butyl-4-hydroxyphenyl)-1,3-oxazol-4-yl]ethyl](methylamino)]propanenitrile, 2,6-ditert-butyl-4-{4-[2-(1-piperazinyl)ethyl]-1,3-oxazol-2-yl}phenol hydrochloride, 2,6-ditert-butyl-4-[4-(hydroxymethyl)-1,3-thiazol-2-yl]phenol and 2,6-ditert-butyl-4-{4-[(methylamino)methyl]-1,3-thiazol-2-yl}phenol hydrochloride.

Cancel **Claim 23**.

Claim 24 (currently amended) The method of claim ~~23~~ 22 wherein the disorder is selected from the group consisting of Parkinson's disease, Alzheimer's disease, Huntington's chorea, amyotrophic lateral sclerosis and peripheral neuropathies.

Claim 25 (previously presented) The method of claim 23 wherein the disorder is pain.